## What is claimed is:

- 1. A carbon monolith comprising a robust carbon monolith characterized by a skeleton size of at least 100 nm, and a hierarchical pore structure having macropores and mesopores.
- 2. A carbon monolith in accordance with claim 1 wherein said carbon monolith is characterized by a skeleton size of 100 nm to 20  $\mu m$ .
- 3. A carbon monolith in accordance with claim 2 wherein said carbon monolith is characterized by a skeleton size of 200 nm to 10 μm.
- 4. A carbon monolith in accordance with claim 3 wherein said carbon monolith is characterized by a skeleton size of 400 nm to 1  $\mu$ m.
- 5. A carbon monolith in accordance with claim 1 wherein said macropores are of a size range of  $0.05 \, \mu m$  to  $100 \, \mu m$ .
- 6. A carbon monolith in accordance with claim 5 wherein said macropores are of a size range of  $0.1 \ \mu m$  to  $50 \ \mu m$ .
- 7. A carbon monolith in accordance with claim 6 wherein said macropores are of a size range of 0.8  $\mu m$  to 10  $\mu m$ .
- 8. A carbon monolith in accordance with claim 1 wherein said mesopores are of a size range of 18 Å to 50 nm.
- 9. A carbon monolith in accordance with claim 8 wherein said mesopores are of a size range of 0.5 nm to 40 nm.
- 10. A carbon monolith in accordance with claim 9 wherein said mesopores are of a size range of 5 nm to 30 nm.

- 11. A carbon monolith in accordance with claim 1 wherein said carbon monolith further comprises graphite.
- 12. A monolithic chromatography column comprising a robust monolithic carbon stationary phase disposed in a chromatography column, said monolithic carbon stationary phase characterized by a skeleton size of at least 100 nm.
- 13. A monolithic chromatography column in accordance with claim 12 wherein said robust monolithic carbon stationary phase is characterized by a skeleton size of 100 nm to 20  $\mu$ m.
- 14. A monolithic chromatography column in accordance with claim 13 wherein said robust monolithic carbon stationary phase is characterized by a skeleton size of 200 nm to 10 μm.
- 15. A monolithic chromatography column in accordance with claim 14 wherein said robust monolithic carbon stationary phase is characterized by a skeleton size of 400 nm to 1 μm.
- 16. A monolithic chromatography column in accordance with claim 12 wherein said monolithic carbon stationary phase is characterized by a hierarchical porous structure.
- 17. A monolithic chromatography column in accordance with claim 16 wherein said hierarchical porous structure comprises macropores and mesopores.
- 18. A monolithic chromatography column in accordance with claim 17 wherein said macropores are of a size range of 0.05 μm to 100 μm.
- 19. A monolithic chromatography column in accordance with claim 18 wherein said macropores are of a size range of  $0.1~\mu m$  to  $50~\mu m$ .
- 20. A monolithic chromatography column in accordance with claim 19 wherein said macropores are of a size range of 0.8  $\mu$ m to 10  $\mu$ m.
- 21. A monolithic chromatography column in accordance with claim 17 wherein said mesopores are of a size range of 18 Å to 50 nm.

- 22. A monolithic chromatography column in accordance with claim 21 wherein said mesopores are of a size range of 0.5 nm to 40 nm.
- 23. A monolithic chromatography column in accordance with claim 22 wherein said mesopores are of a size range of 5 nm to 30 nm.
- 24. A monolithic chromatography column in accordance with claim 12 wherein said monolithic carbon stationary phase further comprises graphite.
- 25. A method of preparing a robust carbon monolith comprising the steps of:
  - a. providing a carbon monolith precursor having a porosity-generating fugitive phase dispersed therein, said fugitive phase comprising mesoparticles and microparticles;
  - b. carbonizing said carbon monolith precursor to form a carbon monolith: and
  - c. removing said fugitive phase from said carbon monolith to form a robust, porous carbon monolith characterized by a skeleton size of at least 100 nm, and a hierarchical pore structure having macropores and mesopores.
- 26. A method in accordance with claim 25 wherein said carbon monolith precursor further comprises at least one carbonizable polymer.
- 27. A method in accordance with claim 25 wherein said porosity-generating fugitive further comprises a material that is soluble in a solvent that does not harm said porous carbon monolith.
- 28. A method in accordance with claim 25 wherein said porosity-generating fugitive further comprises silica.
- 29. A method in accordance with claim 25 further comprising, after said removing step, an additional step of graphitizing said porous carbon monolith.
- 30. A method in accordance with claim 25 wherein said carbon monolith is characterized by a skeleton size of 100 nm to 20  $\mu$ m.

- 31. A method in accordance with claim 30 wherein said carbon monolith is characterized by a skeleton size of 200 nm to  $10 \mu m$ .
- 32. A method in accordance with claim 31 wherein said carbon monolith is characterized by a skeleton size of 400 nm to 1  $\mu$ m.
- 33. A method in accordance with claim 25 wherein said macropores are of a size range of 0.05  $\mu m$  to 100  $\mu m$ .
- 34. A method in accordance with claim 33 wherein said macropores are of a size range of 0.1  $\mu m$  to  $50 \mu m$ .
- 35. A method in accordance with claim 34 wherein said macropores are of a size range of 0.8 μm to 10 μm.
- 36. A method in accordance with claim 25 wherein said mesopores are of a size range of 18 Å to 50 nm.
- 37. A method in accordance with claim 36 wherein said mesopores are of a size range of 0.5 nm to 40 nm.
- 38. A method in accordance with claim 37 wherein said mesopores are of a size range of 5 nm to 30 nm.
- 39. A method of preparing a robust carbon monolith comprising the steps of:
  - a. providing a carbon monolith precursor having a particulate porosity-generating fugitive phase dispersed therein, said fugitive phase comprising mesoparticles and microparticles; and
  - b. heating said carbon monolith precursor to carbonize said carbon monolith precursor, and to remove said fugitive phase from said carbon monolith, to form a robust, porous carbon monolith characterized by a skeleton size of at least 100 nm, and a hierarchical pore structure having macropores and mesopores.

- 40. A method in accordance with claim 39 wherein said carbon monolith precursor further comprises at least one carbonizable polymer.
- 41. A method in accordance with claim 39 wherein said porosity-generating fugitive further comprises a material that is thermally removable at a temperature that does not decompose said porous carbon monolith.
- 42. A method in accordance with claim 39 wherein said porosity-generating fugitive phase further comprises at least one material selected from the group consisting of surfactants and low-charring polymers.
- 43. A method in accordance with claim 39 further comprising, after said removing step, an additional step of graphitizing said porous carbon monolith.
- 44. A method in accordance with claim 39 wherein said carbon monolith is characterized by a skeleton size of 100 nm to 20  $\mu$ m.
- 45. A method in accordance with claim 44 wherein said carbon monolith is characterized by a skeleton size of 200 nm to  $10 \mu m$ .
- 46. A method in accordance with claim 45 wherein said carbon monolith is characterized by a skeleton size of 400 nm to 1 μm.
- 47. A method in accordance with claim 39 wherein said macropores are of a size range of 0.05  $\mu m$  to 100  $\mu m$ .
- 48. A method in accordance with claim 47 wherein said macropores are of a size range of 0.1  $\mu m$  to  $50 \mu m$ .
- 49. A method in accordance with claim 48 wherein said macropores are of a size range of 0.8  $\mu m$  to 10  $\mu m$ .

- 50. A method in accordance with claim 39 wherein said mesopores are of a size range of 18 Å to 50 nm.
- 51. A method in accordance with claim 50 wherein said mesopores are of a size range of 0.5 nm to 40 nm.
- 52. A method in accordance with claim 51 wherein said mesopores are of a size range of 5 nm to 30 nm.
- 53. A method of preparing a monolithic chromatography column comprising the steps of:
  - a. providing a carbon monolith precursor having a porosity-generating fugitive phase dispersed therein;
  - b. carbonizing said carbon monolith precursor to form a carbon monolith;
  - c. removing said fugitive phase from said carbon monolith to form a robust, porous carbon monolith characterized by a skeleton size of at least 100 nm, and a hierarchical pore structure having macropores and mesopores; and
  - d. encapsulating said porous carbon monolith to form a chromatographic column.
- 54. A method in accordance with claim 53 wherein said carbon monolith precursor further comprises at least one carbonizable polymer.
- 55. A method in accordance with claim 53 wherein said porosity-generating fugitive further comprises a material that is soluble in a solvent that does not harm said porous carbon monolith.
- 56. A method in accordance with claim 53 wherein said porosity-generating fugitive further comprises silica.
- 57. A method in accordance with claim 53 further comprising, after said removing step, an additional step of graphitizing said porous carbon monolith.
- 58. A method in accordance with claim 53 wherein said carbon monolith is characterized by a skeleton size of 100 nm to 20 μm.

- 59. A method in accordance with claim 58 wherein said carbon monolith is characterized by a skeleton size of 200 nm to  $10 \mu m$ .
- 60. A method in accordance with claim 59 wherein said carbon monolith is characterized by a skeleton size of 400 nm to 1  $\mu$ m.
- 61. A method in accordance with claim 53 wherein said macropores are of a size range of 0.05  $\mu m$  to  $100 \ \mu m$ .
- 62. A method in accordance with claim 61 wherein said macropores are of a size range of 0.1 μm to 50 μm.
- 63. A method in accordance with claim 62 wherein said macropores are of a size range of 0.8  $\mu m$  to  $10 \ \mu m$ .
- 64. A method in accordance with claim 53 wherein said mesopores are of a size range of 18 Å to 50 nm.
- 65. A method in accordance with claim 64 wherein said mesopores are of a size range of 0.5 nm to 40 nm.
- 66. A method in accordance with claim 65 wherein said mesopores are of a size range of 5 nm to 30 nm.
- 67. A method of preparing a monolithic chromatography column comprising the steps of:
  - a. providing a carbon monolith precursor having a porosity-generating fugitive phase dispersed therein;
  - b. heating said carbon monolith precursor to carbonize said carbon monolith precursor to form a carbon monolith, and to remove said fugitive phase from said carbon monolith; and
  - c. encapsulating said porous carbon monolith to form a chromatographic column.

- 68. A method in accordance with claim 67 wherein said heating step further comprises graphitizing said porous carbon monolith.
- 69. A method in accordance with claim 67 wherein said carbon monolith precursor further comprises at least one carbonizable polymer.
- 70. A method in accordance with claim 67 wherein said porosity-generating fugitive further comprises a material that is thermally removable at a temperature that does not decompose said porous carbon monolith.
- 71. A method in accordance with claim 67 wherein said porosity-generating fugitive phase further comprises at least one material selected from the group consisting of surfactants and low-charring polymers.
- 72. A method in accordance with claim 67 further comprising, after said removing step, an additional step of graphitizing said porous carbon monolith.
- 73. A method in accordance with claim 67 wherein said carbon monolith is characterized by a skeleton size of 100 nm to 20 μm.
- 74. A method in accordance with claim 73 wherein said carbon monolith is characterized by a skeleton size of 200 nm to  $10 \mu m$ .
- 75. A method in accordance with claim 74 wherein said carbon monolith is characterized by a skeleton size of 400 nm to 1  $\mu$ m.
- 76. A method in accordance with claim 67 wherein said macropores are of a size range of 0.05  $\mu m$  to 100  $\mu m$ .
- 77. A method in accordance with claim 76 wherein said macropores are of a size range of 0.1 μm to 50 μm.

- 78. A method in accordance with claim 77 wherein said macropores are of a size range of 0.8  $\mu m$  to 10  $\mu m$ .
- 79. A method in accordance with claim 67 wherein said mesopores are of a size range of 18 Å to 50 nm.
- 80. A method in accordance with claim 79 wherein said mesopores are of a size range of 0.5 nm to 40 nm.
- 81. A method in accordance with claim 80 wherein said mesopores are of a size range of 5 nm to 30 nm.